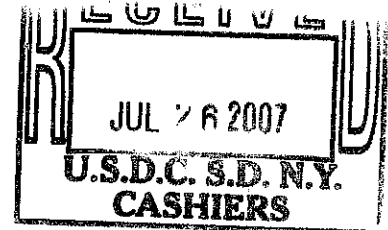


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Attorneys for Plaintiffs



IN THE UNITED STATES DISTRICT COURT  
FOR THE SOUTHERN DISTRICT OF NEW YORK

IN RE: FOSAMAX PRODUCTS LIABILITY  
LITIGATION

MDL No. 1:06-md-1789- JFK

EDITH MACK

Case No. \_\_\_\_\_

Plaintiffs,

v.

**CIVIL COMPLAINT**

MERCK & CO., INC. (hereinafter "Merck"), a  
New Jersey Corporation

**JURY TRIAL DEMANDED**

Defendant.

EDITH MACK, by and through her undersigned counsel, sues Defendant Merck  
& Co., Inc. and allege as follows:

**I. PARTIES**

1. Plaintiff is a resident of the State of North Carolina, and Defendant, Merck, is  
incorporated and has its primary place of business in the State of New Jersey.

2. Plaintiff, EDITH MACK, was born May 4, 1923, and is a resident of Mt. Holly,  
Gaston County, North Carolina. (Unless otherwise specified herein, the term "Plaintiff" refers to

1 Plaintiff, EDITH MACK.) After taking FOSAMAX for an extended period of time, Plaintiff was  
2 diagnosed with serious and permanent injuries.

3 3. Defendant, Merck, is a corporation organized and existing under the laws of the  
4 State of New Jersey, with its principal place of business in New Jersey. The Defendant's  
5 principal office is located at One Merck Drive, Whitehouse Station, New Jersey.  
6

7 4. Defendant, Merck, was at all relevant times authorized to conduct business in the  
8 State of North Carolina.

9 5. Defendant has regularly transacted business in the State of North Carolina and  
10 continues to do so.

11 6. At all relevant times, Defendant, Merck, through its agents, servants, employees,  
12 and apparent agents, was the designer, manufacturer, marketer, distributor, and seller of  
13 FOSAMAX, a bisphosphonate drug used primarily to mitigate or reverse the effects of  
14 osteoporosis, osteopenia, and Paget's Disease.  
15

16 7. Defendant, Merck, either directly or through its agents, apparent agents, servants,  
17 or employees, at all relevant times, sold and distributed FOSAMAX in the State of North  
18 Carolina.  
19

20 8. Defendant encouraged the use of this drug in improper customers, misrepresented  
21 the safety and effectiveness of this drug, and concealed or understated its dangerous side effects  
22 in North Carolina. The Defendant aggressively marketed this drug directly to the consuming  
23 public through the use of various marketing mediums including, but not limited to, print and  
24 television advertisements in North Carolina.

25 9. Based on information and belief, Sales Representatives called physicians on  
26 numerous occasions at which times they presented fraudulent information regarding the safety  
27 and efficacy of FOSAMAX and its harmful side effects, and/or fraudulently suppressed material  
28

1 information regarding the safety and efficacy of FOSAMAX and its harmful side effects, and/or  
2 placed FOSAMAX in the stream of commerce by providing Plaintiff's physician(s) samples of  
3 the drug FOSAMAX.

4  
5 10. At all times material hereto, Merck advertised, marketed, and/or produced  
6 FOSAMAX to Plaintiff utilizing information known to fraudulently represent the safety and  
7 efficacy of FOSAMAX, and said Defendant failed to warn of the known dangers and adverse  
8 events associated with the use of the drug FOSAMAX.

9  
10 11. At all times relevant hereto, the Defendant actually knew of the defective nature of  
11 its product as herein set forth yet continued to design, manufacture, market, distribute, and sell the  
12 product in Gaston County, North Carolina.

13  
14 12. Defendant derives substantial revenue from pharmaceutical products used or  
15 consumed in the State of North Carolina.

16  
17 13. Defendant expected, or should have expected, that its business activities could or  
18 would have consequences within the State of North Carolina.

19  
20 14. Defendant placed FOSAMAX into the stream of worldwide commerce and  
21 interstate commerce in the United States. They did so without adequate testing and with no  
22 warning that the drug carried with it a risk of causing osteonecrosis or osteomyelitis of the jaw.

23  
24 15. Plaintiff needs continued medical monitoring to treat serious and permanent  
25 injuries which have already manifested.

## 26 27 II. JURISDICTION AND VENUE

28  
29 16. This court has jurisdiction pursuant to 28 U.S.C. §§1332, as complete diversity  
30 exists between Plaintiff and Defendant.

31  
32 17. Plaintiff is a resident of the state of North Carolina.

1           18. Defendant, Merck & Co., Inc., is incorporated and has its primary place of  
2 business in the State of New Jersey. The amount in controversy, exclusive of interests and costs,  
3 exceeds \$75,000.00.  
4

5           19. Venue is proper within this district and division pursuant to agreement of the  
6 parties.  
7

### 8                                   **III. FACTUAL BACKGROUND**

9           20. Merck, either directly or through its agents, apparent agents, servants, or  
10 employees, designed, manufactured, marketed, advertised, distributed, and sold FOSAMAX for  
11 the treatment of osteoporosis, Paget's Disease, and other uses.  
12

13           21. As a result of the defective nature of FOSAMAX, persons who were prescribed  
14 and ingested FOSAMAX, including Plaintiff, EDITH MACK, have suffered and may continue to  
15 suffer severe and permanent personal injuries, including osteonecrosis and osteomyelitis.  
16

17           22. Merck concealed and continues to conceal its knowledge of FOSAMAX's  
18 unreasonably dangerous risks from Plaintiff, EDITH MACK, other consumers, and the medical  
19 community.  
20

21           23. Merck failed to conduct adequate and sufficient post-marketing surveillance of  
22 FOSAMAX after it began marketing, advertising, distributing, and selling the drug.  
23

24           24. As a result of Defendant's actions and inaction, Plaintiff, EDITH MACK, was  
25 injured due to her ingestion of FOSAMAX, which has caused and will continue to cause Plaintiff  
26 various injuries and damages. Plaintiff accordingly seeks compensatory damages, as well as  
27 other damages.  
28

          25. At all relevant times, Merck was responsible for, or involved in, designing,  
manufacturing, marketing, advertising, distributing, and selling FOSAMAX.

1           26. In September 1995, the United States Food and Drug Administration ("FDA")  
2 approved Merck's compound alendronate for various uses including the treatment of osteoporosis  
3 and Paget's disease. Defendant, Merck, markets alendronate under the name FOSAMAX.

4           27. FOSAMAX falls within a class of drugs known as bisphosphonates.  
5 Bisphosphonates are used for treating bone conditions such as osteoporosis and Paget's disease.  
6 Other drugs within this class, such as Aredia and Zometa, are used as chemotherapy and as  
7 adjunct chemotherapy but are not indicated for use in non-cancerous conditions such as  
8 osteoporosis.

9  
10           28. There are two classes of bisphosphonates: the N-containing (nitrogenous) and  
11 nonN-containing (non-nitrogenous) bisphosphonates. The nitrogenous bisphosphonates include  
12 the following: pamidronate (Aredia), ibandronate (Bondronat), and alendronate (FOSAMAX).  
13 The non-nitrogenous bisphosphonates include the following: etridonate (Didronel), clodronate  
14 (Bonefos and Loron), and tiludronate (Skelid). Alendronate contains a nitrogen atom. The  
15 Physicians Desk Reference ("PDR") for FOSAMAX confirms that the molecule contains a  
16 nitrogen atom.

17  
18           29. Throughout the 1990s and 2000s, medical articles and studies appeared reporting  
19 the frequent and common occurrence of osteonecrosis of the jaw within the nitrogenous  
20 bisphosphonates used for chemotherapy. As with its reported and acknowledged side effects  
21 concerning irritation, erosion, and inflammation of the upper gastrointestinal tract, Merck knew or  
22 should have known that FOSAMAX, as a nitrogenous bisphosphonate, shared a similar adverse  
23 event profile to the other drugs within this specific subclass of bisphosphonates (i.e., those  
24 containing nitrogen).

25  
26           30. Merck knew, and or should have known, that bisphosphonates, including  
27 FOSAMAX, inhibit endothelial cell function. Similarly, Merck knew, or should have known,  
28

1 that bisphosphonates also inhibit vascularization of the affected area and induce ischemic changes  
2 specific to patients' mandibles (lower jaws) and maxillae (upper jaws) and that these ischemic  
3 changes appear to be cumulative in nature.

4 31. Merck also knew, or should have known, that these factors combine to create a  
5 compromised vascular supply in the affected area. As a result, a minor injury or disease can turn  
6 into a non-healing wound. That, in turn, can progress to widespread necrosis (bone death) and  
7 osteomyelitis (infection of the bone).

8 32. Dentists are now being advised by dental associations to refrain from using any  
9 invasive procedure (such as drilling a cavity) for any patient on FOSAMAX.

10 33. Once the osteonecrosis begins and becomes symptomatic, it is very difficult to  
11 treat and typically is not reversible.

12 34. Shortly after Merck began selling FOSAMAX, reports of osteonecrosis of the jaw  
13 and other dental complications among users began surfacing, indicating that FOSAMAX shared  
14 the class effects of the other nitrogenous bisphosphonates. Despite this knowledge, Merck failed  
15 to implement further study regarding the risk of osteonecrosis of the jaw relative to FOSAMAX.  
16 Rather than evaluating and verifying the safety of FOSAMAX with respect to osteonecrosis of  
17 the jaw, Defendant proposed further uses of FOSAMAX, such as FOSAMAX-D, and sought to  
18 extend the exclusivity period of FOSAMAX through 2018.

19 35. Osteonecrosis of the jaw is a serious medical event and can result in severe  
20 disability and death.

21 36. Since FOSAMAX was released, the FDA has received a significant number of  
22 reports of osteonecrosis of the jaw among users of FOSAMAX.

23 37. On August 25, 2004, the United States Food & Drug Administration ("FDA")  
24 posted its ODS Postmarketing Safety Review on bisphosphonates - - specifically, pamidronate  
25  
26  
27  
28

1 (Aredia), zoledronic acid (Zometa), risedronate (Actonel), and alendronate (FOSAMAX).

2 38. As a result of the FDA Review, the FDA observed that the risk of osteonecrosis of  
3 the jaw was not confined to bisphosphonates used for chemotherapy. The FDA's review  
4 indicated that the osteonecrosis of the jaw was a class effect which specifically extended to the  
5 oral bisphosphonate, FOSAMAX.

6 39. As a result, the FDA recommended and stated that the labeling for FOSAMAX  
7 should be amended by Defendant, Merck, to specifically warn about the risk of osteonecrosis of  
8 the jaw. Defendant, Merck, has refused to accede to the FDA's request and to this day still does  
9 not warn of the risk of osteonecrosis of the jaw in its FOSAMAX labeling.

10 40. Rather than warn patients, and despite knowledge known by Defendant about  
11 increased risk of osteonecrosis of the jaw in patients using FOSAMAX, Defendant continues to  
12 defend FOSAMAX, mislead physicians and the public, and minimize unfavorable findings.

13 41. FOSAMAX is one of Merck's top selling drugs, averaging more than \$3 billion a  
14 year in sales.

15 42. Consumers, including Plaintiff, EDITH MACK, who has used FOSAMAX for the  
16 prevention of osteoporosis, have several alternative, safer products available to treat the  
17 conditions.

18 43. Defendant knew of the significant risk of dental and oral complications caused by  
19 ingestion of FOSAMAX, but Defendant did not adequately and sufficiently warn consumers,  
20 including Plaintiff, EDITH MACK, or the medical community of such risks.

21 44. In an elaborate and sophisticated manner, Defendant aggressively marketed  
22 FOSAMAX directly to consumers and medical professionals (including physicians and leading  
23 medical scholars) in order to leverage pressure on third party payers, medical care organizations,  
24 and large institutional buyers (e.g., hospitals) to include FOSAMAX on their formularies. Faced  
25  
26  
27  
28



1 with the increased demand for the drug by consumers and health care professionals that resulted  
2 from Defendant's successful advertising and marketing blitz, third party payers were compelled  
3 to add FOSAMAX to their formularies. Defendant's marketing campaign specifically targeted  
4 third party payers, physicians, and consumers and was designed to convince them of both the  
5 therapeutic and economic value of FOSAMAX.  
6

7 45. As a direct result, Plaintiff, EDITH MACK, was prescribed FOSAMAX and has  
8 been permanently and severely injured, having suffered serious consequences from the ingestion  
9 of FOSAMAX. Plaintiff, EDITH MACK, requires and will in the future require ongoing medical  
10 care and treatment.

11 46. Plaintiff, EDITH MACK, has suffered from mental anguish from the knowledge  
12 that Plaintiff will have life-long complications as a result of the injuries Plaintiff sustained from  
13 the Plaintiff's use of FOSAMAX.  
14

15 47. Plaintiff, EDITH MACK, was prescribed and began taking FOSAMAX  
16 November 24, 2001.

17 48. Plaintiff used FOSAMAX as prescribed and in a foreseeable manner.

18 49. As a direct and proximate result of using FOSAMAX, Plaintiff suffered  
19 development of serious and permanent injuries..  
20

21 50. Plaintiff, as a direct and proximate result of using FOSAMAX, suffered severe  
22 mental and physical pain and suffering and has sustained permanent injuries and emotional  
23 distress.

24 51. Plaintiff used FOSAMAX which had been provided to her in a condition that was  
25 substantially the same as the condition in which it was manufactured and sold.

26 52. Plaintiff would not have used FOSAMAX had Defendant properly disclosed the  
27 risks associated with the drug.  
28



1           53. Merck, through its affirmative misrepresentations and omissions, actively  
2 concealed from Plaintiff and her physicians the true and significant risks associated with taking  
3 FOSAMAX. The running of any applicable statute of limitations has been tolled by reason of  
4 Merck's fraudulent concealment.

5  
6           54. As a result of Defendant's actions, Plaintiff and her prescribing physicians were  
7 unaware and could not have reasonably known or have learned through reasonable diligence that  
8 Plaintiff had been exposed to the risks identified in this complaint and that those risks were the  
9 direct and proximate result of Defendant's acts, omissions, and misrepresentations.

10           **IV. EQUITABLE TOLLING OF APPLICABLE STATUTES OF LIMITATIONS**

11           55. The running of any statute of limitations has been tolled by reason of Defendant's  
12 fraudulent concealment. Defendant, through its affirmative misrepresentation and omissions,  
13 actively concealed from Plaintiff and her prescribing physician the true risks associated with  
14 taking FOSAMAX.

15  
16           56. As a result of Defendant's actions, Plaintiff and, upon information and belief, her  
17 prescribing physician were unaware, and could not reasonably know or have learned through  
18 reasonable diligence, that she had been exposed to the risks alleged herein and that those risks  
19 were the direct and proximate result of Defendant's acts and omissions.

20  
21           57. Furthermore, Defendant is estopped from relying on any statute of limitations  
22 because of their fraudulent concealment of the true character, quality, and nature of FOSAMAX.  
23 Defendant was under a duty to disclose the true character, quality, and nature of FOSAMAX  
24 because this was non-public information over which the Defendant had and continues to have  
25 exclusive control and because the Defendant knew that this information was not available to the  
26 plaintiffs, medical providers, and/or to their facilities. In addition, the Defendant is estopped  
27 from relying on any statute of limitations because of their intentional concealment of these  
28

1 facts.

2 58. The Plaintiff had no knowledge that the Defendant was engaged in the wrongdoing  
3 alleged herein. Because of the fraudulent acts of concealment of wrongdoing by the Defendant,  
4 the Plaintiff could not have reasonably discovered the wrongdoing at any time prior. Also, the  
5 economics of this fraud should be considered. The Defendant had the ability to and did spend  
6 enormous amounts of money in furtherance of their purpose of marketing and promoting a  
7 profitable drug, notwithstanding the known or reasonably known risks. Plaintiff and medical  
8 professionals could not have afforded and could not have possibly conducted studies to determine  
9 the nature, extent, and identity of related health risks and were forced to rely on only the  
10 Defendant's representations.  
11

## 12 COUNTS

### 13 COUNT I: NEGLIGENCE

14 59 Plaintiff restates the allegations set forth above as if fully rewritten herein.

15 60. The Defendant owed Plaintiff, EDITH MACK, and other consumers a duty to  
16 exercise reasonable care when designing, manufacturing, marketing, advertising, distributing, and  
17 selling FOSAMAX.  
18

19 61. The Defendant failed to exercise due care under the circumstances and therefore  
20 breached this duty by:  
21

- 22 a. Failing to properly and thoroughly test FOSAMAX before releasing the drug  
23 to market;
- 24 b. Failing to properly and thoroughly analyze the data resulting from the pre-  
25 marketing tests of FOSAMAX.
- 26 c. Failing to conduct sufficient post-market testing and surveillance of  
27 FOSAMAX;  
28

- 1 d. Designing, manufacturing, marketing, advertising, distributing, and selling  
2 FOSAMAX to consumers, including Plaintiff, without an adequate warning of  
3 the significant and dangerous risks of FOSAMAX and without proper  
4 instructions to avoid the harm which could foreseeably occur as a result of  
5 using the drug;  
6  
7 e. Failing to exercise due care when advertising and promoting FOSAMAX; and  
8  
9 f. Negligently continuing to manufacture, market, advertise, and distribute  
10 FOSAMAX after Defendant knew or should have known of its adverse effects.  
11  
12 g. Defendant know, or should have known, that consumers, including Plaintiff,  
13 would suffer injuries as a result of Defendant's failure to exercise ordinary  
14 care.

14 62. As a direct and proximate consequence of Defendant's actions, negligence,  
15 omissions, and misrepresentations, Plaintiff, EDITH MACK, has sustained serious and permanent  
16 injuries and will continue to suffer injury, harm, and economic loss.

17 63 Defendant's conduct as described above was committed with knowing, conscious,  
18 wanton, willful, and deliberate disregard for the value of human life and the rights and safety of  
19 consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish  
20 Defendant and deter Defendant from similar conduct in the future.

21  
22 WHEREFORE, Plaintiff demands judgment against Defendant for compensatory and  
23 punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as  
24 the Court deems proper.

25 **COUNT II: STRICT LIABILITY**

26 64 Plaintiff restates the allegations set forth above as if fully rewritten herein.

27 65 Merck manufactured, sold, distributed, marketed, and/or supplied FOSAMAX in a  
28

1 defective and unreasonably dangerous condition to consumers, including Plaintiff, EDITH  
2 MACK. As such, Defendant had a duty to warn the using public, including Plaintiff, of the health  
3 risks associated with using the product.

4 66. Merck designed, manufactured, sold, distributed, supplied, marketed, and/or  
5 promoted FOSAMAX, which was expected to reach and did in fact reach consumers, including  
6 Plaintiff, without substantial change in the condition in which it was manufactured and sold by  
7 the Defendant.

8  
9 67. Plaintiff used FOSAMAX as prescribed and in a manner normally intended,  
10 recommended, promoted, and marketed by Defendant.

11 68. FOSAMAX failed to perform safely when used by ordinary consumers, including  
12 Plaintiff, including when it was used as intended and in a reasonably foreseeable manner.

13 69. FOSAMAX was defective in its design and was unreasonably dangerous in that its  
14 unforeseeable risks exceeded the benefits associated with its design or formulation.

15 70. FOSAMAX was defective in design or formation in that it posed a greater  
16 likelihood of injury than other similar medications and was more dangerous than an ordinary  
17 consumer could reasonably foresee or anticipate.

18 71. FOSAMAX was defective in its design and was unreasonably dangerous in that it  
19 neither bore nor was packaged with nor accompanied by warning adequate to alert consumers,  
20 including Plaintiff, of the risks described herein, including, but not limited to, the risk of  
21 osteonecrosis of the jaw.

22 72. Although Defendant knew, or should have known, of the defective nature of  
23 FOSAMAX, Merck continued to design, manufacture, market, and sell FOSAMAX so as to  
24 maximize sales and profits at the expense of the public health and safety. By so acting, Merck  
25 acted with conscious and deliberate disregard of the foreseeable harm caused by FOSAMAX.  
26  
27  
28

1           73. Plaintiff could not, through the exercise of reasonable care, have discovered  
2 FOSAMAX's defects or perceived the dangers posed by the drug. Plaintiff would not have used  
3 FOSAMAX had the Defendant properly disclosed the risk associated with the drug.

4           74. As a direct and proximate consequence of Defendant's actions, negligence,  
5 omissions, and misrepresentations, Plaintiff, EDITH MACK, has sustained serious and permanent  
6 injuries and will continue to suffer injury, harm, and economic loss.

7           75. Defendant's conduct as described above was committed with knowing, conscious,  
8 wanton, willful, and deliberate disregard for the value of human life and the rights and safety of  
9 consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish  
10 Defendant and deter Defendant from similar conduct in the future.

11           WHEREFORE, Plaintiff demands judgment against Defendant for compensatory and  
12 punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as  
13 the Court deems proper.

14           **COUNT III: BREACH OF EXPRESS WARRANTY**

15           76. Plaintiff restates the allegations set forth above as if fully rewritten herein.

16           77. Defendant expressly represented to Plaintiff, EDITH MACK, and other consumers  
17 and the medical community that FOSAMAX was safe and fit for its intended purposes- that it  
18 was of merchantable quality, that it did not produce any dangerous side effects, and that it was  
19 adequately tested.

20           78. FOSAMAX does not conform to Defendant's express representations because it is  
21 not safe, has numerous side effects, and causes severe and permanent injuries.

22           79. At all relevant times, FOSAMAX did not perform as safely as an ordinary  
23 consumer would expect when used as intended or in a reasonably foreseeable manner.

24           80. Plaintiff, EDITH MACK, other consumers, and the medical community relied  
25  
26  
27  
28

1 upon Defendant's express warranties.

2 81. As a direct and proximate consequence of Defendant's actions, negligence,  
3 omissions, and misrepresentations, Plaintiff, EDITH MACK, has sustained serious and permanent  
4 injuries and will continue to suffer injury, harm, and economic loss.

5 82 Defendant's conduct as described above was committed with knowing, conscious,  
6 wanton, willful, and deliberate disregard for the value of human life and the rights and safety of  
7 consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish  
8 Defendant and deter Defendant from similar conduct in the future.

9 WHEREFORE, Plaintiff demands judgment against Defendant for compensatory and  
10 punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as  
11 the Court deems proper.  
12

13 **COUNT IV: BREACH OF IMPLIED WARRANTY**  
14

15 83 Plaintiff restates the allegations set forth above as if fully rewritten herein.

16 84 Defendant manufactured, distributed, advertised, promoted, and sold FOSAMAX.

17 85 At all relevant times, Defendant knew of the use for which FOSAMAX was  
18 intended and impliedly warranted the product to be of merchantable quality and safe and fit for  
19 such use.  
20

21 86 Merck was aware that consumers, including Plaintiff, EDITH MACK, would use  
22 FOSAMAX for treatment of osteoporosis and for other purposes.

23 87. Plaintiff and the medical community reasonably relied upon the judgment and  
24 sensibility of Merck to sell FOSAMAX only if it was indeed of merchantable quality and safe and  
25 fit for its intended use.

26 88. Defendant breached its implied warranty to consumers, including Plaintiff;  
27 FOSAMAX was not of merchantable quality or safe and fit for its intended use.  
28

1 89. Consumers, including Plaintiff, and the medical community reasonably relied upon  
2 Defendant's implied warranty of FOSAMAX.

3 90. FOSAMAX reached consumers without substantial change in the condition in  
4 which it was manufactured and sold by Defendant.

5 91. As a direct and proximate consequence of Defendant's actions, negligence,  
6 omissions, and misrepresentations, Plaintiff, EDITH MACK, has sustained serious and permanent  
7 injuries and will continue to suffer injury, harm, and economic loss.

8 92. Defendant's conduct as described above was committed with knowing, conscious,  
9 wanton, willful, and deliberate disregard for the value of human life and the rights and safety of  
10 consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish  
11 Defendant and deter it from similar conduct in the future.

12 WHEREFORE, Plaintiff demands judgment against Defendant for compensatory and  
13 punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as  
14 the Court deems proper.

15 **COUNT V: FRAUDULENT MISREPRESENTATIONS**

16 93. Plaintiff restates the allegations set forth above as if fully rewritten herein.

17 94. Merck made fraudulent misrepresentations with respect to FOSAMAX in the  
18 following particulars:

- 19 a. Defendant represented through its labeling, advertising, marketing, materials,  
20 detail persons, seminar presentations, publications, notice letters, and  
21 regulatory submissions that FOSAMAX had been tested and found to be safe  
22 and effective for the prevention and treatment of osteoporosis; and  
23 b. Defendant represented that FOSAMAX was safer than other alternative  
24 medications.  
25  
26  
27  
28



1           95. Defendant knew that their representations were false, yet they willfully, wantonly,  
2 and recklessly disregarded their obligation to provide truthful representations regarding the safety  
3 and risk of FOSAMAX to consumers, including Plaintiff, and the medical community.

4           96. The representations were made by Defendant with the intent that doctors and  
5 patients, including Plaintiff, rely upon them.

6           97. Defendant's representations were made with the intent of defrauding and  
7 deceiving Plaintiff, other consumers, and the medical community to induce and encourage the  
8 sale of FOSAMAX.

9           98. Plaintiff's doctors and others relied upon the representations.

10           99. Defendant's fraudulent representations evinced its callous, reckless, willful, and  
11 depraved indifference to the health, safety, and welfare of consumers, including Plaintiff.

12           100. As a direct and proximate consequence of Defendant's actions, negligence,  
13 omissions, and misrepresentations, Plaintiff, EDITH MACK, has sustained serious and permanent  
14 injuries and will continue to suffer injury, harm, and economic loss.

15           101. Defendant's conduct as described above was committed with knowing, conscious,  
16 wanton, willful, and deliberate disregard for the value of human life and the rights and safety of  
17 consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish  
18 Defendant and deter it from similar conduct in the future.

19           WHEREFORE, Plaintiff demands judgment against Defendant for compensatory and  
20 punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as  
21 the Court deems proper.

22                           **COUNT VI: FRAUDULENT CONCEALMENT**

23           102. Plaintiff restates the allegations set forth above as if fully rewritten herein.

24           103. Merck's fraudulently concealed information with respect to FOSAMAX in the  
25

1 following particulars:

- 2 a. Merck represented through its labeling, advertising, marketing, materials, detail  
3 persons, seminar presentations, publications, notice letters, and regulatory  
4 submissions that FOSAMAX was safe and fraudulently withheld and concealed  
5 information about the substantial risks of using FOSAMAX; and  
6  
7 b. Merck represented that FOSAMAX was safer than other alternative medications  
8 and fraudulently concealed information which demonstrated that FOSAMAX was  
9 not safer than alternatives available on the market.

10 104. Merck had sole access to material facts concerning the dangers and unreasonable  
11 risks of FOSAMAX.

12 105. The concealment of information by Defendant about the risks of FOSAMAX was  
13 intentional, and the representations made by Defendant were known by Defendant to be false.  
14

15 106. The concealment of information and the misrepresentations about FOSAMAX  
16 were made by Defendant with the intent that doctors and patients, including Plaintiff, rely upon  
17 them.

18 107. Plaintiff's doctors and others relied upon the representations and were unaware of  
19 the substantial dental and oral risks of FOSAMAX which Defendant concealed from Plaintiff's  
20 doctors and Plaintiff.

21 108. As a direct and proximate consequence of Defendant's actions, negligence,  
22 omissions, and misrepresentations, Plaintiff, EDITH MACK, has sustained serious and permanent  
23 injuries and will continue to suffer injury, harm, and economic loss.  
24

25 109. Defendant's conduct as described above was committed with knowing, conscious,  
26 wanton, willful, and deliberate disregard for the value of human life and the rights and safety of  
27 consumers such as Plaintiff, thereby entitling Plaintiff to punitive damages so as to punish  
28

1 Defendant and deter it from similar conduct in the future.

2 WHEREFORE, Plaintiff demands judgment against Defendant for compensatory and punitive  
3 damages, together with interest, costs of suit, attorneys' fees, and all such other relief as the Court  
4 deems proper.

5  
6 **COUNT VII: PUNITIVE DAMAGES**

7 110. Plaintiff restates the allegations set forth above as if fully rewritten herein.

8 111. Merck has repeatedly engaged in a pattern of conduct of deliberately avoiding  
9 FDA recommendations relating to public hazards about which the public should be warned.

10 112. For instance, in March 2000, Merck completed a study called VIGOR (VIOXX  
11 Gastrointestinal Outcomes Research) relating to its prescription cox-2 inhibitor, VIOXX. The  
12 VIGOR study showed that VIOXX patients had more than double the rate of serious  
13 cardiovascular problems than those on Naproxen, an older non-steroidal anti-inflammatory drug.  
14 The study was published in the *New England Journal of Medicine*.

15  
16 113. In September 2001, the FDA warned Merck to stop misleading doctors about  
17 VIOXX's effect on the cardiovascular system. Defendant, Merck, was admonished to stop  
18 minimizing the risks of the drug in its marketing. Despite that, Defendant, Merck, refused to  
19 adequately warn physicians and patients about the risk of heart attacks while taking VIOXX.

20  
21 114. On August 25, 2004, a representative from the FDA presented results of a database  
22 analysis of 1.4 million patients. The analysis demonstrated that VIOXX users were more likely to  
23 suffer a heart attack or sudden cardiac death than those taking older non-steroidal drugs. The  
24 FDA representatives concluded that VIOXX was linked to more than 27,000 heart attacks or  
25 sudden cardiac deaths nationwide from the time it came on the market in 1999 through 2003.

26 115. On August 26, 2004, Merck released a press statement which refuted the FDA  
27 analysis and restated Merck's support for the cardiovascular safety of VIOXX.  
28

1           116. On September 30, 2004, Merck recalled VIOXX from the market after having to  
2 halt the APPROVe study (Adenomatous Polyp Prevention on Vioxx). The study was underway  
3 to evaluate the use of VIOXX for recurrent colon polyps. The researchers found an alarming  
4 number of cardiovascular events among the drug users in the APPROVe study.

5           117. At the same time, Merck was aware that the FDA, as of August 24, 2004, was  
6 advising Merck to warn about the risk of osteonecrosis of the jaw for its FOSAMAX patients.  
7 Because Merck knew that its blockbuster drug VIOXX was about to be pulled from the market,  
8 placing more importance on the \$3 billion annual sales of FOSAMAX, Merck deliberately chose  
9 to not amend its packaging of FOSAMAX to include the risk of osteonecrosis of the jaw, fearing  
10 that such a warning would result in reduced revenues for its second largest income producer,  
11 FOSAMAX.  
12

13           118. Merck's acts were willful and malicious in that Merck's conduct was carried on  
14 with a conscious disregard for the safety and rights of Plaintiffs. Defendant's unconscionable  
15 conduct thereby warrants an assessment of exemplary and punitive damages against Merck in an  
16 amount appropriate to punish Merck and deter similar conduct in the future.  
17

18           119. Although Defendant knew or recklessly disregarded the fact that the subject  
19 product causes debilitating and potentially lethal side effects, Defendant continued to market the  
20 subject product to consumers, including Plaintiff, without disclosing these side effects.  
21

22           120. Defendant knew of the subject product's defective nature, as set forth herein, but  
23 continued to design, manufacture, market, and sell it so as to maximize sales and profits at the  
24 expense of the health and safety of the public, including Plaintiff, in conscious and/or negligent  
25 disregard of the foreseeable harm caused by the subject product.

26           121. Defendant intentionally concealed or recklessly failed to disclose to the public,  
27 including Plaintiff, the potentially life-threatening side effects of the subject product to ensure  
28

1 their continued and increased sales. This intentional and/or reckless failure to disclose  
2 information deprived Plaintiff of the information necessary for her to weigh the true risks of using  
3 the subject product against the benefits.

4 122. Defendant's aforementioned conduct was committed with knowing, conscious,  
5 and deliberate disregard for the rights and safety of consumers such as Plaintiff, thereby entitling  
6 Plaintiff to punitive damages in an amount appropriate to punish Defendant and deter it from  
7 similar conduct in the future.

8  
9 WHEREFORE, Plaintiff demands judgment against Defendant for compensatory and  
10 punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as  
11 the Court deems proper.

12 **VIII. PRODUCTS LIABILITY- FAILURE TO WARN**

13  
14 123. Plaintiff restates the allegations set forth above as if fully rewritten herein.

15 124. Defendant designed, tested, manufactured, marketed, sold, and/or distributed  
16 FOSAMAX. As such, it had a duty to warn the using public, including Plaintiff, of the health  
17 risks associated with using the subject product.

18 125. The subject product was under the exclusive control of Defendant and was  
19 unaccompanied by appropriate warnings regarding the health risks associated with its use,  
20 including osteonecrosis of the jaw. The warnings given did not accurately reflect the risk,  
21 incidence, symptoms, scope, or severity of such injury to the consumer. The promotional  
22 activities of Defendant further diluted or minimized the warnings given with the product.

23  
24 126. The subject product was defective and unreasonably dangerous when it left the  
25 possession of the Defendant in that it contained warnings insufficient to alert Plaintiff to the  
26 dangerous risks and reactions associated with it, including, but not limited to, osteonecrosis of the  
27 jaw. Even though Defendant knew or should have known of the risks and reactions associated  
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1 with the subject product, it still failed to provide warnings that accurately reflected the signs,  
2 incidence, scope, or severity of these risks.

3 127. Plaintiff used the subject product for its intended purpose, i.e. for the prevention or  
4 treatment of osteoporosis.

5 128. Plaintiff could not have discovered any defect in the subject product through the  
6 exercise of reasonable care.

7 129. Defendant, as a manufacturer of pharmaceutical drugs, is held to the level of  
8 knowledge of an expert in the field, and further, Defendant had knowledge of the dangerous risks  
9 and side effects of the subject product.

10 130. Plaintiff did not have the same knowledge as Defendant, and no adequate warning  
11 was communicated to her.

12 131. Defendant had a continuing duty to warn consumers, including Plaintiff, of the  
13 dangers associated with the subject product. By negligently and/or wantonly failing to adequately  
14 warn of the dangers of use of the subject product, Defendant breached its duty.

15 132. Although Defendant knew of the defective nature of the subject product, they  
16 continued to design, manufacture, market, and sell it without knowing providing accurate,  
17 adequate, and complete warnings concerning its use so as to maximize sales and profits at the  
18 expense of the public health and safety, in knowing, conscious, and deliberate disregard of the  
19 foreseeable harm caused by the subject product.

20 133. As a direct and proximate result of the Defendant's failure to adequately warn or  
21 other wrongdoing and actions of Defendant described herein, Plaintiff has sustained serious and  
22 permanent injuries, and will continue to suffer, harm, and economic loss.

23 WHEREFORE, Plaintiff demands judgment against Defendant for compensatory and  
24 punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as  
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1 the Court deems proper.

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4 **PRAYER FOR RELIEF**

5 WHEREFORE, Plaintiffs pray for judgment against Defendant as follows:

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1. General damages in an amount to be proven at the time of trial;
  2. Special damages in an amount to be proven at the time of trial;
  3. Exemplary and punitive damages in an amount to be proven at the time of trial  
and sufficient to punish Defendant or to deter Defendant and others from  
repeating the injurious conduct alleged herein;
  4. Pre-judgment and post-judgment interest on the above general and special  
damages;
  5. For costs of this suit and attorney's fees;
  6. All other relief to which Plaintiff may be entitled;
  7. That the costs of this action be taxed to Defendant;
  8. That Plaintiff be granted reasonable attorneys' fees and costs as provided by  
law; and
  9. For such other and further relief as the Court may deem just and proper.



1 Dated: 7/24/2007

REICH & BINSTOCK, LLP

2  
3 By: 

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28 Attorneys for Plaintiffs

**DEMAND FOR JURY TRIAL**

Plaintiff demands a trial by jury on all claims so triable in this action.

Dated: 7/24/2007

REICH & BINSTOCK, LLP

By: 

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